

What is claimed is:

1. An isolated and purified peptide comprising the GD  
5 domain.

2. An isolated and purified peptide having an amino  
acid sequence selected from the group consisting of

GDDINRRYDSEFQ,

PSSTMGQVGRQLAIIGDDINRRYDSEFQ,

QVGRQLAIIGDDINRRYDSEFQTMLQHLQPT,

LSECLKRIGDELDSN,

LKRIGDELD,

QDASTKKLSECLKRIGDELDSNMELQ,

LALRLACIGDEMDVS,

IGDEM,

CMEGSDALALRLACIGDEMDVSLRAPRL,

VGRQLAIIGDDINRR,

and functional equivalents thereof.

3. A mutant of a protein selected from the group  
consisting of Bak, Bax and Bipla, characterized in that it  
comprises the GD domain and exhibits Rat-1 cell killing  
activity substantially equivalent to that of wild-type Bak.

4. A mutant of a protein selected from the group  
consisting of Bak, Bax and Bipla, characterized in that it  
comprises the GD domain and exhibits Bcl-x<sub>L</sub> binding  
substantially equivalent to that of wild-type Bak.



13. The method according to claim 12, wherein said host cell is a mammalian cell.

14. An antibody raised against a GD domain peptide.

15. The antibody of claim 14, wherein said antibody is selected from the group consisting of a polyclonal antibody and a monoclonal antibody.

16. The antibody of claim 15, wherein said antibody is detectably labeled.

17. The antibody of claim 16, wherein said detectable label is selected from the group consisting of: a radio label, an enzyme label, a co-factor label, a fluorescent label, a paramagnetic label, a chemiluminescent label, and a metal label.

18. A detectably labeled nucleotide probe, comprising a first nucleotide sequence which is substantially complementary to a second nucleotide sequence that encodes the GD domain peptide.

19. A pharmaceutical composition comprising a GD domain peptide and a pharmaceutically acceptable carrier.

20. A method of identifying an agent capable of modulating GD domain mediated heterodimerization, comprising:

carrying out a heterodimerization assay which includes a first and a second protein or polypeptide comprising the GD domain, wherein said first and second protein or polypeptide are different, and an agent;

determining whether said agent inhibits or augments heterodimerization of said first protein or polypeptide to said second protein or polypeptide;

wherein if inhibition or augmentation of heterodimerization is determined, it indicates that said agent is capable of modulating GD domain mediated heterodimerization.

21. The method of claim 20, wherein said first and said  
5 second protein or polypeptide is selected from the group consisting of Bak, Bcl-x<sub>L</sub>, Bax and Bipla.

22. A method of identifying an agent capable of modulating GD domain mediated homodimerization, comprising:

10 carrying out a homodimerization assay which includes a first and a second protein or polypeptide comprising the GD domain, wherein said first and second protein or polypeptide are the same, and an agent;

15 determining whether said agent inhibits or augments homodimerization of said first protein or polypeptide to said second protein or polypeptide; wherein if inhibition or augmentation of homodimerization is determined, it indicates that said agent is capable of modulating GD domain mediated homodimerization.

20 23. The method of claim 22, wherein said first and second protein or polypeptide is selected from the group consisting of Bak, Bcl-x<sub>L</sub>, Bax and Bipla.

24. An agent identified by the method of any of claims 20-23.

25 25. Use of an antibody against a GD domain peptide to screen a cDNA expression library for clones comprising DNA inserts encoding immunocrossreactive proteins.

26. An agent comprising a Bcl-2/Bcl-x<sub>L</sub> mimetic.

27. A peptide comprising the GD domain selected from the group consisting of QVG, PEM and derivatives of either of them.